CLAIM AMENDMENTS

1-87 (canceled)

- 88. (new) A method of immunotherapy comprising administering to a patient in need thereof a combination of an alkaloid and a toll-like receptor ligand at a dose sufficient to induce IL-2 production in dendritic cells in the patient.
- 89. (new) The method of claim 88 wherein the immunotherapy comprises:
 - (a) increasing the Th1:Th2 response ratio;
 - (b) haemorestoration;
 - (c) haemoablative immunotherapy;
 - (d) the treatment of immunosuppression;
 - (e) treatment of proliferative disorders (e.g. cancer or cancer metastasis);
 - (f) vaccination, wherein the alkaloid acts as an adjuvant;
 - (g) vaccination, wherein the alkaloid acts to potentiate dendritic cells in situ;
 - (h) wound healing; or
 - (i) the treatment or prophylaxis of infection.
- 90. (new) The method of claim 88 wherein the alkaloid is a piperidine, pyrroline, pyrrolidine, pyrrolizidine, indolizidine or nortropane alkaloid.
- 91. (new) The method of claim 89 wherein the alkaloid is a piperidine, pyrroline, pyrrolizidine, indolizidine or nortropane alkaloid.
- 92. (new) The method of claim 88 wherein the alkaloid is polyhydroxylated.
- 93. (new) The method of claim 89 wherein the alkaloid is polyhydroxylated.

94. (new) The method of claim 92 wherein the alkaloid has the formula:

wherein R is selected from the group comprising hydrogen, straight or branched, unsubstituted or substituted, saturated or unsaturated acyl, alkyl (e.g. cycloalkyl), alkenyl, alkynyl and aryl groups, or a pharmaceutically acceptable salt or derivative thereof.

95. (new) The method of claim 93 wherein the alkaloid has the formula:

$$RO \longrightarrow H \longrightarrow OH$$
 CH_2OH

wherein R is selected from the group comprising hydrogen, straight or branched, unsubstituted or substituted, saturated or unsaturated acyl, alkyl (e.g. cycloalkyl), alkenyl, alkynyl and aryl groups, or a pharmaceutically acceptable salt or derivative thereof.

96. (new) A live cell vaccine comprising an alkaloid and dendritic cells.

97. (new) The vaccine of claim 96 wherein the dendritic cells are antigen-pulsed dendritic cells.

98. (new) The vaccine of claim 96 further comprising T cells.

99. (new) The vaccine of claim 97 further comprising T cells.

100. (new) The vaccine of claim 98 wherein the T cells are primed by contact with dendritic cells.

101. (new) The vaccine of claim 99 wherein the T cells are primed by contact with dendritic cells.

102. (new) The vaccine of claim 100 wherein the T cells are primed by contact with antigen-pulsed dendritic cells.

103. (new) The vaccine of claim 101 wherein the T cells are primed by contact with antigen-pulsed dendritic cells.

104. (new) The vaccine of claim 103 wherein the alkaloid is a piperidine, pyrroline, pyrrolidine, pyrrolizidine, indolizidine or nortropane alkaloid.

105. (new) The vaccine of claim 96 wherein the alkaloid is polyhydroxylated.

106. (new) The vaccine of claim 105 wherein the alkaloid has the formula:

wherein R is selected from the group comprising hydrogen, straight or branched, unsubstituted or substituted, saturated or unsaturated acyl, alkyl (e.g. cycloalkyl), alkenyl, alkynyl and aryl groups, or a pharmaceutically acceptable salt or derivative thereof.

107. (new) A vaccine comprising a neoantigen, an alkaloid and a toll-like receptor ligand.